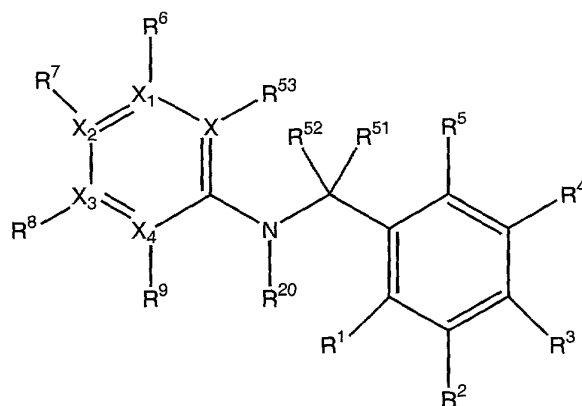


## ABSTRACT

The present invention provides novel compounds of Formula I:



Formula I

its prodrug forms, or pharmaceutically acceptable salts thereof. The compounds of  
 10 this invention are inhibitors of serine proteases, Urokinase (uPA), Factor Xa (FXa),  
 and/or Factor VIIa (FVIIa), and have utility as anti cancer agents and/or as  
 anticoagulants for the treatment or prevention of thromboembolic disorders in  
 mammals. The present invention also provides a process for the selective acylation of  
 an amino group.

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